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PASSWORD:
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                   Zentralblatt
 NEWS 3 OCT 19 BEILSTEIN updated with new compounds
 NEWS 4 NOV 15
                  Derwent Indian patent publication number format enhanced
 NEWS 5 NOV 19 WPIX enhanced with XML display format
NEWS 6 NOV 30 ICSD reloaded with enhancements
NEWS 7 DEC 04 LINPADOCDB now available on STN
 NEWS 8 DEC 14 BEILSTEIN pricing structure to change
 NEWS
          DEC 17 USPATOLD added to additional database clusters
 NEWS 10 DEC 17 IMSDRUGCONF removed from database clusters and STN
NEWS 11 DEC 17 DGENE now includes more than 10 million sequences NEWS 12 DEC 17 TOXCENTER enhanced with 2008 MeSH vocabulary in
                   MEDLINE segment
 NEWS 13 DEC 17 MEDLINE and LMEDLINE updated with 2008 MeSH vocabulary
NEWS 14 DEC 17 CA/CAplus enhanced with new custom IPC display formats
 NEWS 15 DEC 17 STN Viewer enhanced with full-text patent content
                   from USPATOLD
 NEWS 16 JAN 02
                  STN pricing information for 2008 now available
 NEWS 17 JAN 16 CAS patent coverage enhanced to include exemplified
                   prophetic substances
 NEWS 18 JAN 28 USPATFULL, USPAT2, and USPATOLD enhanced with new
                   custom IPC display formats
 NEWS 19 JAN 28 MARPAT searching enhanced
NEWS 20 JAN 28 USGENE now provides USPTO sequence data within 3 days
                   of publication
 NEWS 21 JAN 28 TOXCENTER enhanced with reloaded MEDLINE segment
NEWS 22 JAN 28 MEDLINE and LMEDLINE reloaded with enhancements NEWS 23 FEB 08 STN Express, Version 8.3, now available
                   STN Express, Version 8.3, now available
 NEWS 24 FEB 20 PCI now available as a replacement to DPCI
NEWS 25 FEB 25 IFIREF reloaded with enhancements
NEWS 26 FEB 25 IMSPRODUCT reloaded with enhancements
 NEWS 27 FEB 29 WPINDEX/WPIDS/WPIX enhanced with ECLA and current
                   U.S. National Patent Classification
 NEWS EXPRESS FEBRUARY 08 CURRENT WINDOWS VERSION IS V8.3,
              AND CURRENT DISCOVER FILE IS DATED 20 FEBRUARY 2008
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T.1 STRUCTURE UPLOADED

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SAMPLE SEARCH INITIATED 15:50:12 FILE 'REGISTRY' SAMPLE SCREEN SEARCH COMPLETED -25 TO ITERATE

100.0% PROCESSED 25 ITERATIONS 2 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*
BATCH \*\*COMPLETE\*\* 200 TO PROJECTED ITERATIONS: 800 124 2 TO PROJECTED ANSWERS:

2 SEA SSS SAM L1

=> d 12

ANSWER 1 OF 2 REGISTRY COPYRIGHT 2008 ACS on STN

RN

721959-65-5 REGISTRY Entered STN: 04 Aug 2004 ED

CN  $\beta$ -D-Glucopyranoside, 3-[(3-fluoro-4-methylphenyl)methyl]-4,6-dimethyl-2-pyridinyl (9CI) (CA INDEX NAME)

FS STEREOSEARCH

C21 H26 F N O6 MF

SR

STN Files: CA, CAPLUS, USPATFULL LC

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> s l1 sss full

FULL SEARCH INITIATED 15:50:44 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 593 TO ITERATE

100.0% PROCESSED 593 ITERATIONS 38 ANSWERS

SEARCH TIME: 00.00.01

L3 38 SEA SSS FUL L1

=> file caplus

COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 180.82 181.24

FILE 'CAPLUS' ENTERED AT 15:50:49 ON 27 MAR 2008
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=> s 13

T.4 3 T.3

=> d bib abs hitstr 1-3 14

L4 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2004:996200 CAPLUS

DN 141:424380

TI Preparation of pyrimidine monosaccharides as sodium-dependent glucose transporters and antidiabetic agents

```
ΤN
     Zenkoh, Tatsuya; Eikyu, Yoshiteru; Furukawa, Takako; Kodama, Hiroshi
     Fujisawa Pharmaceutical Co., Ltd., Japan
     PCT Int. Appl., 40 pp.
SO
     CODEN: PIXXD2
DT
     Patent
TιA
     English
FAN.CNT 1
     PATENT NO.
                             KIND
                                     DATE
                                                   APPLICATION NO.
                                                                              DATE
     WO 2004099230
                              A1
                                     20041118
                                                   WO 2004-JP6357
          W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
               CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
               GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
               LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
               NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
               TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
          RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE,
               SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,
               SN, TD, TG
PRAI AU 2003-902263
                                     20030512
    MARPAT 141:424380
OS
GT
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$$\mathbb{R}^4$$
 O O  $\mathbb{R}^1$ 

Pyrimidine monosaccharides I, wherein R1 = H, lower alkyl; R2 = H, lower alkoxy; R3 = H, OH; R4 = H, lower alkoxycarbonyl, hydroxymethyl; and the delocalized bond represents single bond or double bond, or a pharmaceutically acceptable salt thereof, were prepared Pyrimidine monosaccharides I and a salt thereof of the present invention are SGLTs, especially SGIT II selective, inhibitors and are useful for the prevention and/or treatment of diabetes, diabetic retinopathy, diabetic neuropathy, diabetic nephropathy, wound healing, insulin resistance, hyperglycemia, hyper-insulinemia, Syndrome X, diabetic complications, or elevated blood levels of free fatty acids or glycerol, hyperlipidemia, obesity, hypertriglyceridemia, atherosclerosis, hypertension, or for increasing high d. lipoprotein levels and the like, which comprises administering to a mammalian species in need of treatment a therapeutically effective amount of title compds.. Thus, (2S, 3R, 4S, 5R, 6R) - 2 - [[5 - (4 - ethyl - benzyl) - 2 - methoxy - benzyl)]4-pyrimidinyl]oxy]-6-(hydroxymethyl)tetrahydro-2H-pyran-3,4,5-triol was prepared and tested for human SGLT I (IC50 > 100  $\mu M$ ) and SGLT II (IC50 = $0.82~\mu\mathrm{M}$ ) inhibiting activities. In general, amts. between 0.01 mg/ body and about 1,000 mg/kg may be administered per day. 794521-34-9P 794521-35-0P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of pyrimidine monosaccharides as sodium-dependent glucose transporters and antidiabetic agents)

RN 794521-34-9 CAPLUS CN  $\beta$ -D-Glucopyranoside

 $\beta\text{-D-Glucopyranoside, }5\text{-[(4-ethylphenyl)methyl]-2-methoxy-4-pyrimidinyl (9CI) (CA INDEX NAME)}$ 

RN 794521-35-0 CAPLUS

CN β-D-Galactopyranoside, 5-[(4-ethylphenyl)methyl]-2-methoxy-4pyrimidinyl (9CI) (CA INDEX NAME)

Absolute stereochemistry.

IT 794521-29-2P 794521-30-5P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of pyrimidine monosaccharides as sodium-dependent glucose transporters and antidiabetic agents)

RN 794521-29-2 CAPLUS

β-D-Glucopyranoside, 5-[(4-ethylphenyl)methyl]-2-methoxy-4-pyrimidinyl, 2,3,4,6-tetrabenzoate (9CI) (CA INDEX NAME)

 ${\tt Absolute \ stereochemistry.}$ 

RN 794521-30-5 CAPLUS

CN  $\beta$ -D-Galactopyranoside, 5-[(4-ethylphenyl)methyl]-2-methoxy-4-pyrimidinyl, 2,3,4,6-tetrabenzoate (9CI) (CA INDEX NAME)

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

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ANSWER 2 OF 3 CAPLUS COPYRIGHT 2008 ACS on STN
L4
       2004:566633 CAPLUS
ΑN
DN
       141:123854
      Preparation of D-glucose derivatives as human SGLT2 inhibitors Fujikura, Hideki; Nishimura, Toshihiro; Katsuno, Kenji; Isaji, Masayuki Kissei Pharmaceutical Co., Ltd., Japan
TΙ
IN
PA
       PCT Int. Appl., 75 pp. CODEN: PIXXD2
SO
DT
       Patent
LA
       Japanese
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My Application - tcm

FAN.CNT 1																			
							)	DATE			APPLICATION NO.						DATE		
PI	WO	70 2004058790					A1 200407			WO 2003-JP16310						20031219			
		W:	ΑE,	AG,	AL,	AM,	ΑT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BW,	BY,	BZ,	CA,	CH,	
			CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,	
			GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KR,	KΖ,	LC,	LK,	
			LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	ΝI,	NO,	NΖ,	
			OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	ΤJ,	TM,	
			TN,	TR,	TΤ,	TΖ,	UA,	UG,	US,	UΖ,	VC,	VN,	YU,	ZA,	ZM,	ZW			
		RW:	BW,	GH,	GM,	KE,	LS,	MW,	MΖ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	ΑZ,	
			BY,	KG,	KΖ,	MD,	RU,	ТJ,	TM,	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	
			ES,	FΙ,	FR,	GB,	GR,	HU,	IE,	ΙT,	LU,	MC,	NL,	PT,	RO,	SE,	SI,	SK,	
			TR,	BF,	ΒJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	ΤG
	CA	2509835									CA 2003-2509835								
	ΑU	U 2003289440					A1 20040722				AU 2	003-	2894	40		20031219			
	EΡ	> 1577317				A1 20050921			EP 2003-780923						20031219				
		R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙT,	LΙ,	LU,	NL,	SE,	MC,	PT,	
			IE,	SI,	LT,	LV,	FΙ,	RO,	MK,	CY,	AL,	TR,	BG,	CZ,	EE,	HU,	SK		
		S 2006035840							US 2005-540519					20050623					
PRAI	JΡ	JP 2002-374016					A 20021225												
	WO 2003-JP16310							2003	1219										
OS	MAF	RPAT	141:	1238	54														

AB The title compds. I [wherein X1-X4 = independently N, (un)substituted CH, etc.; R  $\neq$  4-Z-Ph; Z = H, halo, (un)substituted alkyl, alkoxy, etc.]

ΙI

McIntosh

GI

Absolute stereochemistry.

RN 721959-66-6 CAPLUS CN  $\beta$ -D-Glucopyranoside, 3-[(3-methylphenyl)methyl]-2-pyridinyl (9CI) (CA INDEX NAME)

Absolute stereochemistry.

IT 721959-70-2P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(intermediate; preparation of D-glucose derivs. as human SGLT2 inhibitors)

RN 721959-70-2 CAPLUS

CN  $\beta$ -D-Glucopyranoside, 3-[(3-fluoro-4-methylphenyl)methyl]-4,6-dimethyl-2-pyridinyl, 2,3,4,6-tetraacetate (9CI) (CA INDEX NAME)

Absolute stereochemistry.

L4 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2003:5979 CAPLUS

DN 138:49945

TI Nitrogenous heterocyclic derivative, medicinal composition containing the same, medicinal use thereof, and intermediate therefor

IN Nishimura, Toshihiro; Fujikura, Hideki; Fushimi, Nobuhiko; Tatani, Kazuya; Katsuno, Kenji; Isaji, Masavuki

Katsuno, Kenji; Isaji, Masayuki PA Kissei Pharmaceutical Co., Ltd., Japan

SO PCT Int. Appl., 102 pp.

CODEN: PIXXD2

DT Patent

LA Japanese

FAN.CNT

E AIV .		_																		
	PATENT NO.						D	DATE		APPLICATION NO.						DATE				
						-														
PI	WO	2003	<b>A</b> 1	A1 20030103			1	WO 2	002-	JP60	20020617									
		W:	ΑE,	AG,	AL,	AM,	ΑT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	BZ,	CA,	CH,	CN,		
			co,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,		
			GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KR,	KΖ,	LC,	LK,	LR,	LS,		
			LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	NZ,	OM,	PH,	PL,		
			PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ΤJ,	TM,	TN,	TR,	TT,	ΤZ,	UA,		
			UG,	US,	UΖ,	VN,	YU,	ZA,	ZM,	ZW										
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			CY,	DE,	DK,	ES,	FI,	FR,	GB,	GR,	IE,	ΙT,	LU,	MC,	NL,	PT,	SE,	TR,		
			BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	ΤG		
	CA	2455300				A1		2003	0103	CA 2002-2455300						20020617				
	ΑU	AU 2002313248					A1 20030108				AU 2002-313248						20020617			
	ΕP	EP 1405859						2004	0407		EP 2	002-	7387:		20020617					

these compounds are excluded via a proviso in my app-tcm

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R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
                                      20041013
                                                    CN 2002-814975
      CN 1537114
                                                                                 20020617
                               А
      BR 2002010510
                               Α
                                      20050111
                                                    BR 2002-10510
                                                                                 20020617
      US 2005049203
                               Α1
                                      20050303
                                                    US 2004-481013
                                                                                 20040820
      US 7271153
                               В2
                                      20070918
PRAI JP 2001-187368
                                      20010620
                               Α
     WO 2002-JP6000
                               T<sub>0</sub>T
                                      20020617
OS
      MARPAT 138:49945
GΙ
```

AΒ A nitrogenous heterocyclic derivative represented by the general formula (I), a pharmacol. acceptable salt thereof, or a prodrug of either. These have excellent human SGLT2 inhibitory activity and are useful as a preventive or remedy for diseases attributable to hyperglycemia such as diabetes. In the general formula [I; X1 and X3 each is nitrogen or CH; X2 is nitrogen or CR2; X4 is nitrogen or CR3 (provided that one or two of X1 to X4 are nitrogen); and R1, R2, and R3 are hydrogen, etc.]. 479481-37-3P,  $2-(2,3,4,6-Tetra-O-acetyl-\beta-D$ glucopyranosyloxy) -3-[4-(2-Methoxymethyloxyethyl) benzyl]-4,6dimethylpyridine 479481-38-4P RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses) (preparation of nitrogenous heterocyclic derivs. as antidiabetics and for other medicinal uses) 479481-37-3 CAPLUS RN CN  $\beta$ -D-Glucopyranoside, 3-[[4-[2-(methoxymethoxy)ethyl]phenyl]methyl]-4,6-dimethyl-2-pyridinyl, 2,3,4,6-tetraacetate (9CI) (CA INDEX NAME)

Ι

Absolute stereochemistry.

RN 479481-38-4 CAPLUS CN  $\beta$ -D-Glucopyranoside, 3-[[4-[2-(methoxymethoxy)ethyl]phenyl]methyl]-4,6-dimethyl-2-pyridinyl (9CI) (CA INDEX NAME)

479481-29-3P,  $6-(N-Acetylamino)-2-(2,3,4,6-tetra-0-acetyl-\beta-D-1)$ glucopyranosyloxy)-3-(4-ethylbenzyl)pyridine 479481-31-7P, 6-Amino-2-(2,3,4,6-tetra-0-acetyl- $\beta$ -D-glucopyranosyloxy)-3-(4ethylbenzyl)pyridine 479481-33-9P 479481-35-1P,  $2-(2,3,4,6-\text{Tetra-O-acetyl-}\beta-\text{D-glucopyranosyloxy})-3-(4-\text{methoxybenzyl})-3-(4-\text{meth$ 4,6-dimethylpyridine 479481-41-9P 479481-43-1P 479481-45-3P 479481-46-4P, 4-(2,3,4,6-Tetra-O-acetyl- $\beta$ -D-glucopyranosyloxy)-5-(4-methoxybenzyl)-2,6-dimethylpyridine 479481-48-6P 479481-50-0P, 4-(2,3,4,6-Tetra-O-acetyl- $\beta$ -D-glucopyranosyloxy) -5-(4-Ethylbenzyl)-2,6-dimethylpyrimidine 479481-52-2P, 4-(2,3,4,6-Tetra-O-acetyl-β-D- $\verb|glucopyranosyloxy|| -3 - (4 - \verb|butylbenzyl||) -2, 6 - \verb|dimethylpyridine||$ 479481-54-4P 479481-56-6P, 3-(2,3,4,6-Tetra-O-acetyl- $\beta$ -D-glucopyranosyloxy)-4-benzylpyridazine 479481-57-7P 479481-58-8P 479481-59-9P 479481-60-2P 479481-61-3P,  $2-(\beta-D-Glucopyranosyloxy)-3-[4-(2-\beta-D-Glucopyranosyloxy)]$ hydroxyethyl)benzyl)-4,6-dimethylpyridine 479481-62-4P 479481-63-5P,  $4-(4-Ethoxybenzyl)-3-(\beta-D-$ Glucopyranosyloxy) pyridine 479481-64-6P, 2-( $\beta$ -D- ${\tt Glucopyranosyloxy)-3-(4-methoxybenzyl)\,pyridine}\quad {\tt 479481-65-7P,}$  $4-\left(\beta-D-\texttt{Glucopyranosyloxy}\right)-5-\left(4-\texttt{methoxybenzyl}\right)-2,6-\texttt{dimethylpyrimidine}$ 479481-66-8P,  $3-(\beta-D-Glucopyranosyloxy)-4-[4-(2$ hydroxyethyl)benzyl)pyridine 479481-67-9P, 5-(4-Ethylbenzyl)-4- $(\beta-D-Glucopyranosyloxy)-2,6-dimethylpyrimidine 479481-68-0P$ ,  $3-(4-Butylbenzyl)-4-(\beta-D-Glucopyranosyloxy)-2,6-dimethylpyridine$ 479481-69-1P 479481-70-4P 479481-71-5P RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of nitrogenous heterocyclic derivs. as antidiabetics and for other medicinal uses) 479481-29-3 CAPLUS RN Acetamide, N-[5-[(4-ethylphenyl)methyl]-6-[(2,3,4,6-tetra-O-acetyl- $\beta$ -D-glucopyranosyl)oxy]-2-pyridinyl]- (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-31-7 CAPLUS
CN β-D-Glucopyranoside, 6-amino-3-[(4-ethylphenyl)methyl]-2-pyridinyl,
2,3,4,6-tetraacetate (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-33-9 CAPLUS CN  $\beta$ -D-Glucopyranoside, 3-[(4-ethylphenyl)methyl]-4,6-dimethyl-2-pyridinyl, 2,3,4,6-tetraacetate (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-35-1 CAPLUS CN  $\beta$ -D-Glucopyranoside, 3-[(4-methoxyphenyl)methyl]-4,6-dimethyl-2-pyridinyl, 2,3,4,6-tetraacetate (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-41-9 CAPLUS CN  $\beta$ -D-Glucopyranoside, 6-methoxy-3-[(4-methoxyphenyl)methyl]-4-methyl-2-pyridinyl, 2,3,4,6-tetraacetate (9CI) (CA INDEX NAME)

RN 479481-43-1 CAPLUS CN  $\beta$ -D-Glucopyranoside, 4-[(4-ethoxyphenyl)methyl]-3-pyridinyl, 2,3,4,6-tetraacetate (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-45-3 CAPLUS CN  $\beta$ -D-Glucopyranoside, 3-[(4-methoxyphenyl)methyl]-2-pyridinyl, 2,3,4,6-tetraacetate (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-46-4 CAPLUS
CN β-D-Glucopyranoside, 3-[(4-methoxyphenyl)methyl]-2,6-dimethyl-4pyridinyl, 2,3,4,6-tetraacetate (9CI) (CA INDEX NAME)

RN 479481-48-6 CAPLUS CN  $\beta$ -D-Glucopyranoside, 4-[[4-[2-(benzoyloxy)ethyl]phenyl]methyl]-3-pyridinyl, 2,3,4,6-tetraacetate (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-50-0 CAPLUS CN  $\beta$ -D-Glucopyranoside, 5-[(4-ethylphenyl)methyl]-2,6-dimethyl-4-pyrimidinyl, 2,3,4,6-tetraacetate (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-52-2 CAPLUS CN  $\beta$ -D-Glucopyranoside, 3-[(4-butylphenyl)methyl]-2,6-dimethyl-4-pyridinyl, 2,3,4,6-tetraacetate (9CI) (CA INDEX NAME)

RN 479481-54-4 CAPLUS CN  $\beta$ -D-Glucopyranoside, 3-[(4-methoxyphenyl)methyl]pyrazinyl, 2,3,4,6-tetraacetate (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-56-6 CAPLUS CN  $\beta$ -D-Glucopyranoside, 4-(phenylmethyl)-3-pyridazinyl, 2,3,4,6-tetraacetate (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-57-7 CAPLUS CN Acetamide, N-[5-[(4-ethylphenyl)methyl]-6-( $\beta$ -D-glucopyranosyloxy)-2-pyridinyl]- (CA INDEX NAME)

RN 479481-58-8 CAPLUS CN  $\beta$ -D-Glucopyranoside, 6-amino-3-[(4-ethylphenyl)methyl]-2-pyridinyl (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-59-9 CAPLUS CN  $\beta$ -D-Glucopyranoside, 3-[(4-ethylphenyl)methyl]-4,6-dimethyl-2-pyridinyl (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-60-2 CAPLUS CN  $\beta$ -D-Glucopyranoside, 3-[(4-methoxyphenyl)methyl]-4,6-dimethyl-2-pyridinyl (9CI) (CA INDEX NAME)

RN 479481-61-3 CAPLUS
CN β-D-Glucopyranoside, 3-[[4-(2-hydroxyethyl)phenyl]methyl]-4,6dimethyl-2-pyridinyl (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-62-4 CAPLUS
CN β-D-Glucopyranoside, 6-methoxy-3-[(4-methoxyphenyl)methyl]-4-methyl-2pyridinyl (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-63-5 CAPLUS 
CN  $\beta$ -D-Glucopyranoside, 4-[(4-ethoxyphenyl)methyl]-3-pyridinyl (9CI) (CA INDEX NAME)

RN 479481-64-6 CAPLUS CN  $\beta$ -D-Glucopyranoside, 3-[(4-methoxyphenyl)methyl]-2-pyridinyl (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-65-7 CAPLUS CN  $\beta$ -D-Glucopyranoside, 5-[(4-methoxyphenyl)methyl]-2,6-dimethyl-4-pyrimidinyl (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-66-8 CAPLUS CN  $\beta$ -D-Glucopyranoside, 4-[[4-(2-hydroxyethyl)phenyl]methyl]-3-pyridinyl (9CI) (CA INDEX NAME)

RN 479481-67-9 CAPLUS CN  $\beta$ -D-Glucopyranoside, 5-[(4-ethylphenyl)methyl]-2,6-dimethyl-4-pyrimidinyl (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-68-0 CAPLUS CN  $\beta$ -D-Glucopyranoside, 3-[(4-butylphenyl)methyl]-2,6-dimethyl-4-pyridinyl (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-69-1 CAPLUS CN  $\beta$ -D-Glucopyranoside, 3-[(4-methoxyphenyl)methyl]pyrazinyl (9CI) (CA INDEX NAME)

RN 479481-70-4 CAPLUS CN  $\beta$ -D-Glucopyranoside, 4-(phenylmethyl)-3-pyridazinyl (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 479481-71-5 CAPLUS CN  $\beta$ -D-Glucopyranoside, 3-[(4-methoxyphenyl)methyl]-4,6-dimethyl-2-pyridinyl, 6-(methyl carbonate) (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RE.CNT 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

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(FILE 'HOME' ENTERED AT 15:48:48 ON 27 MAR 2008)

FILE 'REGISTRY' ENTERED AT 15:49:39 ON 27 MAR 2008

L1 STRUCTURE UPLOADED

L2 2 S L1

L3 38 S L1 SSS FULL

FILE 'CAPLUS' ENTERED AT 15:50:49 ON 27 MAR 2008

L4 3 S L3